

05-25-04

JFW 3754

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE Washington, D.C., United States of America

In re Application of HU, X. D.

Serial No.: 09/851,177

Filed: May 8, 2001

For: HIGH SURFACE AREA, SMALL

CRYSTALLITE SIZE CATALYST FOR FISCHER-TROPSCH SYNTHESIS

REQUEST FOR WITHDRAWAL OF HOLDING OF ABANDONMENT BASED ON REPLY TIMELY FILED AND RECEIVED IN THE OFFICE BUT NOT CORRELATED WITH THE APPLICATION PURSUANT TO 37 CFR 1.181

Commissioner for Patents Group Director 1754 P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

We kindly request the United States Patent Office to reinstate patent application 09/851,177 in accordance with 37 CFR § 1.181 based on the following facts:

- 1. We responded to the March 26, 2003 (3/26/03) office action by fax on September 24, 2003 (Exhibit A).
- 2. Included was a Petition for Extension of Time Under 37 CFR 1.136(a) for three months with payment by credit card of \$930.00 (Exhibit B).
- 3. We received the Auto-Reply Facsimile Transmission Report on September 24, 2003 with time as 8:36:34 AM (EDT) (Exhibit C).
- 4. We received a Notice of Abandonment on November 18, 2003 (Exhibit D).
- 5. On November 26, 2003 in accordance with 37 CFR 1.8 (b), we informed the

USPTO we previously transmitted by fax a response to the office action with mailing date of 03/26/03 (Exhibit E).

6. Since that time we have contacted the Inventor's Assistance Center on numerous occasions that informed us in December and January the application was still abandoned. February, March and April it was stated by the Assistance Center that the application was with the Group Art Unit 1754. We left numerous messages with Stan Silverman who is the head of Group 1700. In May, we were told by the Center they could not tell from the information on the computer exactly where this application was located.

Having satisfied all requirements, applicants respectfully request that this Petition be granted and the holding of abandonment of the application be withdrawn.

Further the applicant respectfully request that the USPTO treat the Response to the March 26, 2003 Office Action as timely filed.

Respectfully submitted,

Joan Simunic, Reg. No. 43,125

Süd-Chemie Inc. (502) 634-7373

jsimunic@sud-chemieinc.com

Date: 5/25/04

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The USPTO date stamp acknowledges receipt of the paper(s) listed

Patent/Application No.: 09/851,177 Applicant: Sud-Chemie Inc.

Filing Date: May 8, 2001

Title: HIGH SURFACE AREA, SMALL CRYSTALLITE SIZE CATALYST

FOR FISCHER-TROPSCH SYNTHESIS

Payment: N/A

Attachments: Cover letter; copy of response to office action; copy of auto reply facsimile transmission; copy of notice of

USPS Express Mail #: ER 521694377 US

Response to notice of abandonment





Süd-Chemie Inc. 1600 West Hill Street Louisville, Kentucky 40210

PHONE: (502) 634-7373 FAX: (502) 634-7724

CONFIDENTIAL AND PRIVILEGED ATTORNEY CLIENT COMMUNICATION

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TO:

U.S. Patent and Trademark Office

Examiner: Cam Nguyen

Group: 1754

Fax #703-872-9306

FROM:

Joan L. Simunic Reg. No. 43,125

DATE:

September 24, 2003

PAGES:

20 in total (including cover sheet)

RE:

U.S. Patent Application No.09/851,177

Remarks: This facsimile is in response to the office action, having a mailing date of March 26, 2003

Enclosed are:

- (1) Transmittal Form.
- (2) Fee Transmittal for FY 2003
- (3) Credit Card Payment Form
- (4) Petition for Extension of Time
- (5) Response to Office Action
- (6) Clean Unmarked Version of Claims Now in Application

CERTIFICATE OF FACSIMILE TRANSMISSION UNDER 37 CFR 1.8

I hereby certify that this paper is being facsimile transmitted to the Patent and Trademark Office on the date shown below.

Donna Ferrill

Person Signing

100/ 11/10

9/24/03

Date

PTO/SB/21 (08-03)

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	FORM		First Named Inventor	X. D. Hu			
(to be used for a	all correspondence after initial	filing)	Art Unit	1754			
			Examiner Name	Cam Ngu	ıyen		
Total Number of	Pages in This Submission	20	Attorney Docket Number	ZL 494/0	1001		
		ENCL	OSURES (Check all that	t apply)			
Fee Transmittal Form Drawing(s)							
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Date	September 23, 2008	-					
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sufficient postage a	I hereby certify that this correspondence is being facsimile transmitted to the USPTO or deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450 on the date shown below.						
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This collection of information is required by 37 CFR 1.5. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

App. for use through 07/31/2006. OMB 0651-0032
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Effective 01/01/2003. Patent fees are subject to annual revision.

Applicant claims small entity status. See 37 CFR 1.27

TOTAL AMOUNT OF PAYMENT

Complete if Known		
Application Number	09/851,177	
Filing Date	May 8, 2001	
First Named Inventor	X. D. Hu	
Examiner Name	Cam Nguyen	
Art Unit	1754	
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METHOD OF PAYMENT (check all that apply) FEE CALCULATION (continued)				
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(Complete (if applicable)) SUBMITTED BY Telephone 502-634-7373 Registration No. 43,125 Jean L. Simunic (Attomey/Agent) Name (Print/Type) September 23, 2003 Date Signature

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This collection of information is required by 37 CFR 1.17 and 1.27. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

PTO/SB/22 (08403)

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Ϋ ÉΤ	TION FOR EXTENSION OF TIME UNI	DER 37 CFR 1.136(a) Docket Number (Optional) 71494/0100 (
		In re Application of X.D. Hu et al
		Application Number 09/851, 177 Filed May 8, 2001
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		Art Unit 1734 Examiner C. Navyen
appli	cation.	1.136(a) to extend the period for filing a reply in the above identified
The		nall-entity fee are as follows (check time period desired):
	One month (37 CFR 1.17(a)(1))	\$
	Two months (37 CFR 1.17(a)(2))	\$
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	Four months (37 CFR 1.17(a)(4))	\$
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	I am the applicant/inventor.	
	Statement under 3	the entire interest. See 37 CFR 3.71. 7 CFR 3.73(b) is enclosed (Form PTO/SB/96).
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This collection of information is required by 37 CFR 1.136(a). The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 6 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

PTO/SB/22 (08403)

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PETITION FOR EXTENSION OF TIME UND	ER 37 CFR 1.136(a)	Docket Number (Optional) 71494/0100 (
	In re Application of X.	D. Hu et al
	Application Number 09	/851, 177 Filed May 8, 2001
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Four months (37 CFR 1.17(a)(4))		\$
Five months (37 CFR 1.17(a)(5))		\$
Applicant claims small entity status. See 37 half, and the resulting fee is: \$		ee amount shown above is reduced by one-
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I am the applicant/inventor.		
	he entire interest. See 37 CFR 3.73(b) is enclosed	
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NOTE: Signatures of all the inventors or assignees of record of signature is required, see below.	the entire interest or their represen	ntative(s) are required. Submit multiple forms if more than one
	ns are submitted.	

This collection of information is required by 37 CFR 1.136(a). The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 6 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application Number

09/851,177

Applicants

X. D. Hu et al.

Filing Date

05/08/2001

TC/A.U.

1754

Examiner

Cam Nguyen

Attorney Docket No.

ZL494/01001

Custor Title:

Customer No. :

High Surface Area, Small Crystallite Size Catalyst for Fischer-Tropsch

Synthesis

Commissioner for Patents P.O. Box 1450

Alexandria VA 22313-1450

ADMENDMENT

Sir:

In response to the Office Action mailed March 26, 2003, please amend the above-identified application as follows:

Amendments to the Specification: There are no changes are being made to the specification.

Amendments to the Claims are reflected in the listing of claims which begins on page 2 of this paper.

Amendments to the Drawings: There are no changes are being made to the drawings.

Remarks/Arguments begin on page 6 of this paper.

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1. (currently amended) A catalyst for use in the Fischer-Tropsch process, said catalyst comprising a catalyst particle, which comprises at least one metal that is an efficient carbon monoxide adsorber and at least one promoter, said metal and said promoter being dispersed on a support to form a said catalyst particle, said particle having a BET surface area of from about 100 m²/g to about 250 m²/g, and said metal and said promoter being are dispersed on the support such that the crystallite size of the metal oxide is from about 40 Å to about 200 Å, and said particle having an essentially smooth, homogeneous surface morphology.

Claim 2. (original) The catalyst of Claim 1 wherein said particle comprises from about 5 wt % to about 60 wt % cobalt, and from about 0.0001 wt % to about 1 wt % of a first promoter, and from about 0.01 wt % to about 5 wt % of a second promoter.

Claim 3. (original) The catalyst of Claim 2 wherein said particle comprises from about 10 wt% to about 30 wt % cobalt, and from about 0.01 wt % to about 0.05 wt % of said first promoter, and from about 0.1 wt % to about 1 wt % of said second promoter.

Claim 4. (original) The catalyst of Claim 1 wherein said metal is selected from the group consisting of nickel, cobalt, iron, ruthenium, osmium, platinum, palladium, iridium, rhenium, molybdenum, chromium, tungsten, vanadium, rhodium, copper, zinc, and combinations thereof.

Claim 5. (original) The catalyst of Claim 4 wherein said metal is cobalt.

Claim 6. (currently amended) The catalyst of Claim 1 wherein said promoter is selected from the group consisting of boron, cerium, chromium, copper, iridium, iron, lanthanum, manganese, molybdenum, palladium, platinum, rhenium, rhodium, ruthenium, strontium, tungsten, vanadium, zinc, sodium oxide, potassium oxide, rubidium oxide, cesium oxide, magnesium oxide, titanium oxide, zirconium oxide, and other rare earth metals, such as scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, other rare earth metals and combinations thereof.

Claim 7. (original) The catalyst of Claim 2 wherein said first promoter is selected from the group consisting of palladium, platinum, ruthenium, rhenium, rhodium, iridium and a combination thereof; and said second promoter is selected from the group consisting of potassium, boron, cesium, lanthanum, cerium, strontium, scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, palladium, platinum, ruthenium, rhenium, rhodium, iridium and combinations thereof.

Claim 8. (original) The catalyst of Claim 1 wherein said support is selected from the group consisting of aluminum oxide, γ -alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.

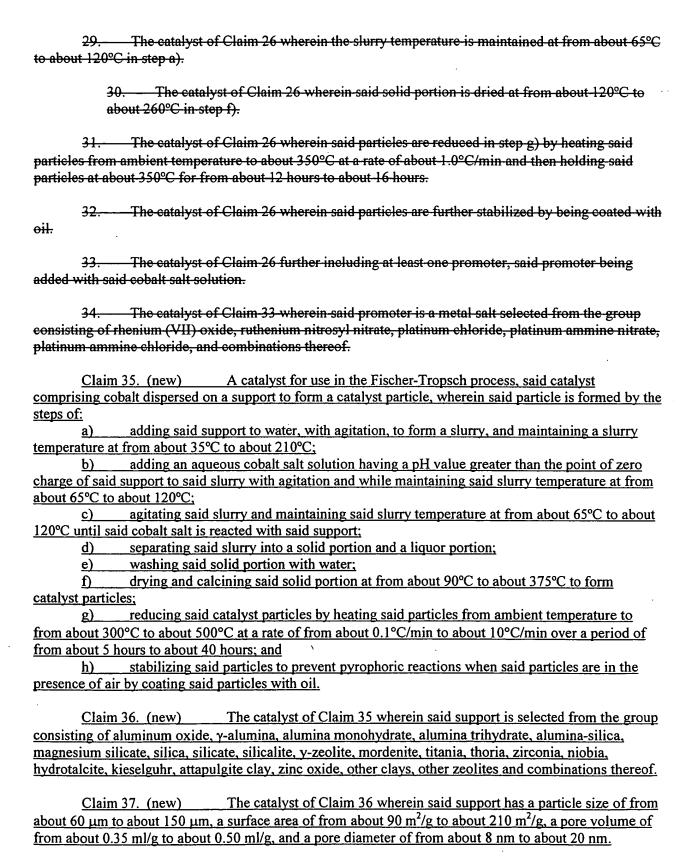
Claim 9. (original) The catalyst of Claim 8 wherein said support is γ -alumina.

- Claim 10. (original) The catalyst of Claim 9 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.
- Claim 11. (currently amended) A catalyst for use in the Fischer-Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size, and wherein said particle being is formed by the steps of:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- c) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is essentially completely reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles; and
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours.
- Claim 12. (original) The catalyst of Claim 11 wherein said support is selected from the group consisting of aluminum oxide, γ -alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silicate, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
 - Claim 13. (original) The catalyst of Claim 12 wherein said support is aluminum oxide.
- Claim 14. (original) The catalyst of Claim 12 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.
- Claim 15. (original) The catalyst of Claim 11 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.
- Claim 16. (currently amended) The catalyst of Claim 15 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water molecules, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.
- Claim 17. (original) The catalyst of Claim 16 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate.
- Claim 18. (original) The catalyst of Claim 11 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).

- Claim 19. (original) The catalyst of Claim 11 wherein said solid portion is dried at from about 120°C to about 260°C in step f).
- Claim 20. (original) The catalyst of Claim 11 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- Claim 21. (original) The catalyst of Claim 11 wherein said particles are further stabilized to prevent pyrophoric reactions when said particles are in the presence of air.
- Claim 22. (currently amended) The catalyst of Claim 21 wherein said particles are stabilized by being coated coating with oil.
- Claim 23. (currently amended) The catalyst of Claim 11 further including at least one promoter, wherein said promoter being is added with said cobalt salt solution.
- Claim 24. (original) The catalyst of Claim 23 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 25. (original) The catalyst of Claim 11 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.

Claims 26 - 34. (canceled)

- 26. A method for making a catalyst for use in the Fischer Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle, said method comprising:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt-salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- e) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is essentially completely reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles; and
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours.
 - 27. The catalyst of Claim 26 wherein said support is aluminum oxide.
- 28. The catalyst of Claim 27 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate.



Claim 38. (new) The catalyst of Claim 35 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.

Claim 39. (new) The catalyst of Claim 38 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.

Claim 40. (new) The catalyst of Claim 35 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).

Claim 41. (new) The catalyst of Claim 35 wherein said solid portion is dried at from about 120°C to about 260°C in step f).

Claim 42. (new) The catalyst of Claim 35 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.

Claim 43. (new) The catalyst of Claim 35 further including at least one promoter wherein said promoter is added with said cobalt salt solution.

Claim 44. (new) The catalyst of Claim 43 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.

Claim 45. (new) The catalyst of Claim 35 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.

Claim 46. (new) The catalyst of Claim 35 wherein said support is aluminum oxide and said aqueous cobalt salt solution comprises water and hexaammine cobalt (II) carbonate.

Attachment: Clean Unmarked Version of Claims Now in Application

REMARKS / ARGUMENTS

Remarks Regarding Informalities and Claims Rejected Under 35 USC §112

Claims 1-25 remain in the application. Claims 35-46 have been added to the application and are presented for examination. Claims 26-34 have been withdrawn in response to a requirement by the Examiner that the claims be restricted to Claims 1-25, drawn to a catalyst, or to Claims 26-34, drawn to a process of preparing a catalyst. In view of the Examiner's earlier restriction requirement, applicant retains the right to present claims 26-34 in a divisional application.

The Examiner objected to claims 1, 11, 16, 22, and 23 because of noted informalities. As proposed by the Examiner, claims 1 and 11 have been reworded in lines 1-3. Further, as proposed by the Examiner: in claim 1, line 5, "being" has been replaced with "are"; in claim 11, line 2, "being" has been replaced with "is"; and, in claim 16, line 2, "molecules" has been deleted to retain consistency with the language of claim 15. Further formality amendments include: in claim 22, line 1, "being coated" has been deleted and replaced with "coating"; and in claim 23, line 2, "being" has been replaced with "is" and "wherein" has been added. The amendments of claim 22 and claim 23 were suggested by the Examiner but required further correction than proposed to have grammatically correct sentences after amendment. Finally, in claim 11, step (c), "essentially completely" has been deleted to address a concern raised by the Examiner with the original language. No new subject matter has been added in making these amendments.

The Examiner objected to claim 17 based on the failure to comply to 37 CFR 1.75(c). The applicant respectfully challenges this objection. Claim 17 depends from claim 16 which recites a cobalt (II) complex having "coordination sphere ligands selected from the group consisting of water, ammonia, ..." Claim 17 recites "The catalyst of Claim 16 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate." As is known in the art, the "hexaammine cobalt (II) carbonate" coordination sphere ligands are ammonia, as recited in claim 16. The carbonate is not a coordination sphere ligand but rather a counterion to the positively charged cobalt ion. Thus, applicant requests the Examiner to reconsider his objection with respect to claim 17.

Claims 1-10 and 16-17 were rejected under 35 U.S.C. §112. The Examiner states that Claim 1 recites the limitation "the metal oxide" with insufficient antecedent basis for this limitation. The applicant respectfully notes that in the specification on page 4, lines 2-3, the catalyst has "a metal oxide crystallite size of from about 40 Å to about 200 Å." On page 12, lines 11-15, the metal oxide is more narrowly defined as the "cobalt oxide crystallite size" – and as is known in the art, cobalt is considered a metal when in reference to oxides – and the "cobalt oxide crystallite size ... is greater than about 40 Å diameter, and is preferably less than about 200 Å." Thus, applicant requests the Examiner to reconsider his objection under 35 U.S.C. §112 with respect to claim 1, and its dependent claims 2-10. If it would expedite matters, the applicant is willing to amend the paragraph on page 12 to include the more inclusive language of page 4 if so requested by the Examiner.

As noted by the Examiner, Claim 6 included the indefinite phrase "such as". Claim 6 has been amended to delete this phrase and should now be in allowable form.

As noted by the Examiner, Claim 16 included improper Markush terminology. Claim 16 has been amended to be consistent with the correct Markush phrase, and claim 16 and its dependent claim 17 should now be in allowable form.

Remarks Regarding Allowable Subject Matter

The Examiner has indicated that Claim 22 is objected to as being dependent on a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims. In response to this rejection, new claim 35 is presented. New claim 35 includes the limitations of rejected independent claims 11 (preamble and steps a - g), rejected dependent claim 21 and objected to dependent claim 22 (step h). Claims 36 - 46 have been added as claims dependent on new independent claim 35. As amended, independent claim 35 is believed to now be in allowable form. Because they depend from an allowable claim, dependent claims 36 - 46 are also believed to now be in allowable form.

The Examiner has also indicated that claims 1-10 would be allowable if the 35 U.S.C. §112 rejection is overcome. As noted in the prior section, page 4, lines 2-3, of the specification teach a catalyst that has "a metal oxide crystallite size of from about 40 Å to about 200 Å" and page 12, lines 11 -15, of the specification teach "cobalt oxide crystallite size ... greater than about 40 Å diameter, and is preferably less than about 200 Å." Applicant is willing to amend the paragraph on page 12 to include the more inclusive language of page 4 if so requested by the Examiner.

Remarks Regarding Citations

Applicant has made note of the prior art recited by the Examiner in Paragraph 15 of his Office Action mailed March 26, 2003.

Remarks Regarding Claims Rejected Under 35 USC §102(b) and 35 USC §103(a)

The Examiner has rejected independent claim 11 and dependent claims 12, 14 – 21, 23 and 25 under 35 U.S.C 102(b) as anticipated by or, in the alternative, under 35 U.S.C. 103(a) as obvious over Shen et al, U.S. Patent 5,962,367 ("the '367 patent"). Applicant respectfully traverses both rejections.

The Examiner has further rejected independent claim 11 and dependent Claims 12 - 13, 15 - 21 and 23 - 25 under 35 U.S.C 102(b) as anticipated by or in the alternative, under 35 U.S.C. 103(a) as obvious over Sapienza et al., U.S. Patent 4,396,539 ("the '539 patent"). Applicant respectfully traverses both rejections.

Summary of the Present Invention

The Fischer-Tropsch catalyst of the present invention is a transition metal-based catalyst having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support, and a small metal oxide crystallite size. In a first embodiment, the catalyst has a surface area of from about 100 m²/g to about 250 m²/g; an essentially smooth, homogeneous surface morphology; an essentially uniform distribution of metal throughout an essentially inert support; and a metal oxide crystallite size of from about 40 Å to about 200 Å. Optionally, the catalyst may further comprise at least one promoter.

The Fischer-Tropsch catalyst of independent claim 11 of the present application, as amended, is a catalyst comprising cobalt dispersed on a support to form a catalyst particle having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size, and wherein the particle is formed by several prescribed steps. Claims 12-25 of the present application depend from independent claim 11.

As provided in dependent claim 14, the catalyst of claim 11 may be prepared from a support having a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm. However, as discussed on page 28, line 1 through page 29, line 4 and as shown in Figures 1A – 4, two catalysts prepared with the same support material can have significantly different physical characteristics depending upon the process used to make the catalyst. (*N.B.*: In the examples of the present application each catalyst was prepared using a Puralox SCCa 5/150 support having a surface area of about 160 m²/g, a pore volume of about 0.50 ml/g, and a pore diameter of about 12.55 nm.) The steps outlined in independent claim 11 result in a catalyst particle having the desired physical characteristics. Prior art methods of catalyst particle preparation, even starting with the same support, do not result in the catalyst particle of the present invention.

Summary of U.S. Patent 5,962,367, Shen et al.

U.S. Patent 5,962,367 teaches a process for preparing a catalyst support primarily formed of titania. The catalyst support comprises 60 wt% to 100 wt % titania as TiO₂ and 0 wt% to 40 wt % alumina as Al₂O₃. As noted by the Examiner, the '367 patent teaches a cobalt molybdate catalyst supported on a titania support, wherein the support has a surface area ranging from 80 m²/g to 200 m²/g, a pore volume of from about 0.3 ml/g to about 0.5 ml/g, and a pore diameter of from about 6 nm to about 20 nm.

The '367 patent then teaches using this catalyst support to prepare a cobalt molybdate catalyst using an impregnation process. At column 5, lines 6 – 8, the '367 states "the dynamic co-impregnation

method is used for supporting the cobalt and molybdenum active components ..." In Examples 4-7, the support is "immersed into the impregnating solution ..." for the preparation of the catalyst.

However, the '367 patent does not teach or suggest that the catalyst resulting from the impregnation process using the '367 titania support has a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size as is seen with the preparation process of the present invention. Rather, there is nothing in the '367 patent that suggests that the physical characteristics of the cobalt molybdate catalyst prepared by the impregnation process would be any different than the catalysts prepared by the prior art impregnation method and reported in Example 6 (and Figures 2A, 2B and 4) of the present application. Further, the catalyst of the present invention made by the process of instant claim 11 would not be anticipated or obvious merely because a titania support is used, or the physical characteristics of the support are similar to supports of the prior art, or because a promoter is added. Thus, independent claim 11 and its dependent claims 12 – 25 are not anticipated nor obvious in view of U.S. Patent 5,962,367.

Summary of U.S. Patent 4,396,539, Sapienza et al.

U.S. Patent 4,396,539 teaches a process for preparing a catalyst suitable for Fischer-Tropsch synthesis wherein the catalyst is composed of palladium or platinum and cobalt supported on a solid support. More specifically, the '539 patent teaches an alumina, silica gel, kieselguhr, or zinc oxide support that is impregnated by immersion of the support in an aqueous solution of the salt of a palladium or platinum metal. The palladium- or platinum-treated support is then immersed in a solution containing primarily cobalt.

The objective of the process set forth in the '539 patent is to achieve a catalyst particle with the metal loading as a coating around the support. The catalyst structure depicted in Figure 5 is believed to "best represent the idealized structure of this catalyst" (column 4, lines 9 – 21). By contrast, the process of claim 11 of the present invention produces a catalyst that has "an essentially uniform distribution of cobalt throughout the support". In other words, the process taught in the '539 patent teaches away from the desired product of the present invention. Thus, independent claim 11 and its dependent claims 12 -25 are not anticipated nor obvious in view of U.S. Patent 4,396,539.

Because U.S. Patent 5,962,367 and U.S. Patent 4,396,539 each teach a process for preparing a catalyst that relies on impregnation to deliver the metal to the support, the product-by-process taught by the amended claim 11 of the present application is neither anticipated nor obvious. Thus, applicant respectfully requests that the §102(b) and the §103(a) rejections be withdrawn and that independent claim 11, as amended, and its dependent claims 12-25, be allowed.

Applicant respectfully requests that a timely Notice of Allowance be issued in this case. Respectfully submitte

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Clean Unmarked Version of Claims Now in Application

- Claim 1. A catalyst for use in the Fischer-Tropsch process, said catalyst comprising a catalyst particle, which comprises at least one metal that is an efficient carbon monoxide adsorber and at least one promoter dispersed on a support to form said catalyst particle, said particle having a BET surface area of from about 100 m²/g to about 250 m²/g, and said metal and said promoter are dispersed on the support such that the crystallite size of the metal oxide is from about 40 Å to about 200 Å, and said particle having an essentially smooth, homogeneous surface morphology.
- Claim 2. The catalyst of Claim 1 wherein said particle comprises from about 5 wt % to about 60 wt % cobalt, and from about 0.0001 wt % to about 1 wt % of a first promoter, and from about 0.01 wt % to about 5 wt % of a second promoter.
- Claim 3. The catalyst of Claim 2 wherein said particle comprises from about 10 wt% to about 30 wt % cobalt, and from about 0.01 wt % to about 0.05 wt % of said first promoter, and from about 0.1 wt % to about 1 wt % of said second promoter.
- Claim 4. The catalyst of Claim 1 wherein said metal is selected from the group consisting of nickel, cobalt, iron, ruthenium, osmium, platinum, palladium, iridium, rhenium, molybdenum, chromium, tungsten, vanadium, rhodium, copper, zinc, and combinations thereof.
 - Claim 5. The catalyst of Claim 4 wherein said metal is cobalt.
- Claim 6. The catalyst of Claim 1 wherein said promoter is selected from the group consisting of boron, cerium, chromium, copper, iridium, iron, lanthanum, manganese, molybdenum, palladium, platinum, rhenium, rhodium, ruthenium, strontium, tungsten, vanadium, zinc, sodium oxide, potassium oxide, rubidium oxide, cesium oxide, magnesium oxide, titanium oxide, zirconium oxide, scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, other rare earth metals and combinations thereof.
- Claim 7. The catalyst of Claim 2 wherein said first promoter is selected from the group consisting of palladium, platinum, ruthenium, rhenium, rhodium, iridium and a combination thereof; and said second promoter is selected from the group consisting of potassium, boron, cesium, lanthanum, cerium, strontium, scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, palladium, platinum, ruthenium, rhenium, rhodium, iridium and combinations thereof.
- Claim 8. The catalyst of Claim 1 wherein said support is selected from the group consisting of aluminum oxide, γ-alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silica, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
 - Claim 9. The catalyst of Claim 8 wherein said support is γ -alumina.
- Claim 10. The catalyst of Claim 9 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.

- Claim 11. A catalyst for use in the Fischer-Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle_having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size, and wherein_said particle is formed by the steps of:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- c) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles; and
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours.
- Claim 12. The catalyst of Claim 11 wherein said support is selected from the group consisting of aluminum oxide, γ -alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silicate, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
 - Claim 13. The catalyst of Claim 12 wherein said support is aluminum oxide.
- Claim 14. The catalyst of Claim 12 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.
- Claim 15. The catalyst of Claim 11 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.
- Claim 16. The catalyst of Claim 15 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.
- Claim 17. The catalyst of Claim 16 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate.
- Claim 18. The catalyst of Claim 11 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).
- Claim 19. The catalyst of Claim 11 wherein said solid portion is dried at from about 120°C to about 260°C in step f).

- Claim 20. The catalyst of Claim 11 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- Claim 21. The catalyst of Claim 11 wherein said particles are further stabilized to prevent pyrophoric reactions when said particles are in the presence of air.
 - Claim 22. The catalyst of Claim 21 wherein said particles are stabilized by coating with oil.
- Claim 23. The catalyst of Claim 11 further including at least one promoter wherein said promoter is added with said cobalt salt solution.
- Claim 24. The catalyst of Claim 23 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 25. The catalyst of Claim 11 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.
- Claim 35. A catalyst for use in the Fischer-Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle, wherein said particle is formed by the steps of:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- c) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles;
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours; and
- h) stabilizing said particles to prevent pyrophoric reactions when said particles are in the presence of air by coating said particles with oil.
- Claim 36. The catalyst of Claim 35 wherein said support is selected from the group consisting of aluminum oxide, γ-alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silica, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
- Claim 37. The catalyst of Claim 36 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.

- Claim 38. The catalyst of Claim 35 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.
- Claim 39. The catalyst of Claim 38 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.
- Claim 40. The catalyst of Claim 35 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).
- Claim 41. The catalyst of Claim 35 wherein said solid portion is dried at from about 120°C to about 260°C in step f).
- Claim 42. The catalyst of Claim 35 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- Claim 43. The catalyst of Claim 35 further including at least one promoter wherein said promoter is added with said cobalt salt solution.
- Claim 44. The catalyst of Claim 43 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 45. The catalyst of Claim 35 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.
- Claim 46. The catalyst of Claim 35 wherein said support is aluminum oxide and said aqueous cobalt salt solution comprises water and hexaammine cobalt (II) carbonate.

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Applicant claims small entity status. See 37 CFR 1.27

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First Named Inventor	X. D. Hu			
Examiner Name	Cam Nguyen			
Art Unit	1754			
Attorney Docket No.	ZL 494/01001			

METHOD OF PAYMENT (check all that apply) FEE CALCULATION (continued)				
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1. BASIC FILING FEE 1253 930 2253 465 Extension for reply within third month Large Entity Small Entity Fee Bold 1253 930 2253 465 Extension for reply within fourth month	-			
Fee Fee Fee Fee Description Fee Paid 1254 1,450 2254 725 Extension for reply wild in local minutes	<u> </u>			
Code (\$) Code (\$) 1255 1,970 2255 985 Extension for reply within fifth month				
1401 320 2401 160 Notice of Appeal				
1402 320 2402 160 Filing a brief in support of an appear				
1403 280 2403 140 Request for oral hearing				
1451 1,510 Petition to institute a public use proceed	ding			
SUBTOTAL (1) (\$) n/a				
1453 1.300 2453 650 Petition to revive - uninternational				
2. EXTRA CLAIM FEES FOR UTILITY AND REISSUE 1501 1,300 2501 650 Utility issue fee (or reissue)	-			
Extra Claims below Fee Paid 1502 470 2502 235 Design issue ree				
Total Claims20** = X = 1503 630 2503 315 Plant issue fee				
Independent Claims 1460 130 1460 130 Petitions to the Commissioner				
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Large Entity Small	Sum -			
Fee Fee Fee Fee Description Code (\$) Code (\$) Code (\$)	' <u>-</u>			
1202 18 2202 9 Claims in excess of 20 1809 750 2809 375 Filling a submission after final rejection				
1301 84 2201 42 Independent claims in excess of 3 (37 CFR 1.129(a))				
1203 280 2203 140 Multiple dependent claim, if not paid 1810 750 2810 375 For each additional invention to be examined (37 CFR 1.129(b))				
1204 84 2204 42 ** Reissue independent daims	(RCE) -			
1802 900 1802 900 Request for expedited examination	_			
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Other fee (specify) -	<u>-</u>			
SUBTOTAL (2) (3) *Reduced by Basic Filing Fee Paid SUBTOTAL (3) (\$	930			

				(Complete (if applicable))
SUBMITTED BY	Jean L. Simunic	Registration No.	43,125	Telephone	502-634-7373
Name (Print/Type)	aball L. Silliume	(Attorney/Agent)		Date	September 23, 2003
Cionatum					

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PET	ITION FOR EXTENSION OF TIME UND	ER 37 CFR 1.136(a)	Docket Number (Optional) 71494/0100 (
		In re Application of X.	D. Hu et al		
		Application Number 09	1851, 177 Filed May 8, 2001		
		For High Surfa	ce Azea		
		Art Unit 1754	Examiner C. Nauyen		
	is a request under the provisions of 37 CFR $^\circ$ cation.	1.136(a) to extend the perio	, · · · .		
The	requested extension and appropriate non-sm	all-entity fee are as follows	(check time period desired):		
	One month (37 CFR 1.17(a)(1))		\$		
	Two months (37 CFR 1.17(a)(2))		\$		
	Three months (37 CFR 1.17(a)(3))		s 930 -		
	Four months (37 CFR 1.17(a)(4))		\$		
	Five months (37 CFR 1.17(a)(5))		\$		
	Applicant claims small entity status. See 37 half, and the resulting fee is: \$	CFR 1.27. Therefore, the fo	ee amount shown above is reduced by one-		
	A check in the amount of the fee is enc	losed.			
X	Payment by credit card. Form PTO-203	88 is attached.			
	The Director has already been authorized to charge fees in this application to a Deposit Account.				
	The Director is hereby authorized to ch to Deposit Account Number	arge any fees which may	be required, or credit any overpayment,		
	I have enclosed a duplicate copy of this	s sheet.			
	I am the applicant/inventor.	·			
		he entire interest. See 37 CFR 3.73(b) is enclosed			
	attorney or agent of red	cord. Registration Numbe	er <u>43,125</u>		
	attorney or agent unde Registration number if a	r 37 CFR 1.34(a). cting under 37 CFR 1.34(a)			
	WARNING: Information on this form may on this form. Provide credit card informa				
	Syptember 23, 200:	3	Signature Signature		
	502-634-7373 Telephone Number		Typed or printed name		
	E: Signatures of all the inventors or assignees of record of ture is required, see below.	the entire interest or their represen	tative(s) are required. Submit multiple forms if more than one		
		ns are submitted.			

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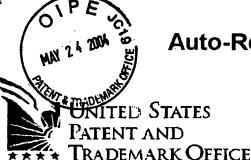
PTO/SB/22 (08403)

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PET	TION	FOR	EXT	ENSION OF TIME UND	ER 37 CFR 1.136(a)	Docket Number (Optional) 71494/0100 (
					In re Application of	χ.	D. Hu et al
					Application Numbe	09	7/851,177 Filed May 8, 2001
				•	For High Sc	rsa	ce Acca
					Art Unit 175	1	Examiner C. Kauyen
	s a requation.	uest (under	the provisions of 37 CFR	1.136(a) to extend the	e perio	od for filing a reply in the above identified
The r	equest	ed ex	tensio	on and appropriate non-sm	all-entity fee are as f	ollows	(check time period desired):
		One	mon	th (37 CFR 1.17(a)(1))			\$
		Two	mon	ths (37 CFR 1.17(a)(2))			\$
	X	Thre	e mo	nths (37 CFR 1.17(a)(3))			s <u>930</u>
		Fou	r mon	ths (37 CFR 1.17(a)(4))			s
		Five	mon	ths (37 CFR 1.17(a)(5))			\$
<u>.</u>				small entity status. See 37 Ilting fee is: \$	CFR 1.27. Therefore	e, the t	fee amount shown above is reduced by one-
	A Little of the factor and and						
X	Payment by credit card. Form PTO-2038 is attached.						
	The Director has already been authorized to charge fees in this application to a Deposit Account.						
	The Director is hereby authorized to charge any fees which may be required, or credit any overpayment, to Deposit Account Number						
	I have enclosed a duplicate copy of this sheet.						
	l am	the		applicant/inventor.			
				assignee of record of the Statement under 3			7 CFR 3.71. ed (Form PTO/SB/96).
			X	attorney or agent of re	cord. Registration	Numb	per 43,125
				attorney or agent under Registration number if a	er 37 CFR 1.34(a). acting under 37 CFR 1.3	4(a) _	
	WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.						
		Sy	ote	Date 23, 200	з 🧲	10	Signature
	-	50	2-	634-7373 Telephone Number	_	17	Typed or printed name
	: Signati ture is re				f the entire interest or their	represe	entative(s) are required. Submit multiple forms if more than one
		al of			ms are submitted.	•	

This collection of information is required by 37 CFR 1.136(a). The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 6 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

TO:Auto-reply fax to 32 634 7661 COMPANY:



Auto-Reply Facsimile Transmission

TO:

Fax Sender at 502 634 7661

Fax Information Date Received:

9/24/03 8:36:34 AM [Eastern Daylight Time]

Total Pages:

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09/24/03 WED 07:47 FAX 802 634 7651

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Süd-Chemie Inc. 1600 West Hill Street Louisville, Kentucky 40210

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TO

U.S. Patent and Trademark Office Examiner: Cam Nguyen Gro Fax **8703-872-9310** Croup: 1754

FROM: Joan L.Simunic Reg. No. 43,125

CATE:

September 24, 2003

PAGES :

30 in total (including cover sheet)

U.S. Patent Application No.09/851,177

Remarks: This facinile is in response to the office action, having a mailing date of March $26,\ 2003$

Enclosed are:

(1) Transmittal form.
(2) For Wransmittal for AV 2003
(3) Credit Card Payment Form
(4) Potinion for Extension of Time
(5) Response to Office Action
(6) Clean Unmarked Version of Claims Now in Application

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UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22113-1450

APPLICATION NO.	FII	LING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/851,177	0	5/08/2001	X.D. Hu	ZL494/01001	2722
22884	7590	11/18/2003		EXAM	INER
MIDDLETON & REUTLINGER 2500 BROWN & WILLIAMSON TOWER				NGUYEN	, CAM N
LOUISVILL	E, KY 4	0202		ART UNIT	PAPER NUMBER

- Notice 2 abendonment

DATE MAILED: 11/18/2003

dic.

RECEIVED NOV 2

Please find below and/or attached an Office communication concerning this application or proceeding.

PTO-90C (Rev. 10/03)



Application No.

Applicant(s) 09/851,177

Cam Nguyen

Examiner

Art Unit

1754

Hu et al.



-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

The state of the s	
This application is abandoned in view of:	
1. X Applicant's failure to timely file a proper reply to the Office letter mailed on Mar 26, 2003	,
(a) A reply was received on (with a Certificate of Mailing or Transmission dated), which is after the expiration of the period for reply (including a total extension of time of month(s)) which expired on	
(b) A proposed reply was received on, but it does not constitute a proper reply under 37 CFR 1.113(a) to the final rejection.	
(A proper reply under 37 CFR 1.113 to a final rejection consists only of: (1) a timely filed amendment which places the application in condition for allowance; (2) a timely filed Notice of Appeal (with appeal fee); or (3) a timely filed Request for Continued Examination (RCE) in compliance with 37 CFR 1.114).	
(c) A reply was received on but it does not constitute a proper reply, or a bona fide attempt at proper reply, to the non-final rejection. See 37 CFR 1.85(a) and 1.111. (See explanation in box 7 below).	а
(d) X No reply has been received.	
2. Applicant's failure to timely pay the required issue fee and publication fee, if applicable, within the statutory period three months from the mailing date of the Notice of Allowance (PTOL-85).	Of
(a) The issue fee and publication fee, if applicable, was received on (with a Certificate of Mailing of Transmission dated), which is after the expiration of the statutory period for payment of the issue fee (and publication fee) set in the Notice of Allowance (PTOL-85).	or Sue
(b) The submitted issue fee of \$ is insufficient. A balance of \$ is due.	
The issue fee required by 37 CFR 1.18 is \$ The publication fee, if required by 37 CFR 1.18(d) is \$	
(c) The issue fee and publication fee, if applicable, has not been received.	_
3. Applicant's failure to timely file corrected drawings as required by, and within the three-month period set in, the Notice of Allowability (PTO-37).	
(a) Proposed new formal drawings were received on (with a Certificate of Mailing or Transmission dated), which is after the expiration of the period for reply.	
(b) ☐ No corrected drawings have been received.	
4. The letter of express abandonment which is signed by the attorney or agent of record, the assignee of the entire interest, or all of the applicants.	
5. The letter of express abandonment which is signed by an attorney or agent (acting in a representative capacity under 37 CFR 1.34(a)) upon the filing of a continuing application.	
6. The decision by the Board of Patent Appeals and Interferences rendered on and because the period for seeking court review of the decision has expired and there are no allowed claims.	
7. The reason(s) below:	
LISAR I	
Petitions to revive under 37 CFR 1.137(a) or (b), or requests to withdraw the holding of abandonment under 37 CFR 1.181 should be promptly filed to minimize any negative effects on patent term.	,

IN THE ENDEED STATES PATENT AND TRADEMA 'K OFFICE Washington, D.C., United States of America

In re Application of HU, X.D.; et al.

Serial No.: 09/851,177

Filed: 05/08/2001

For: HIGH SURFACE AREA, SMALL

CRYSTALLITE SIZE CATALYST FOR

FISCHER-TROPSCH SYNTHESIS

Mail Stop Non-Fee Director of Technology Center Art Unit 1754 P.O. Box 1450 Alexandria, VA 22313-1450

CERTIFICATE OF TRANSMISSION UNDER 37 CFR 1.8 (b)

Dear Sir:

We kindly request the United States Patent Office to consider the enclosed response to the 3/26/03 office action timely filed as it was faxed on September 24, 2003 and can be evidenced by the Auto-Reply Facsimile Transmission report.

In accordance with 37 CFR 1.8 (b), we are promptly informing the Office we previously transmitted by fax a response to the office action with mailing date of 03/26/03. I hereby state the response was sent by fax on 9/24/03 by Donna Ferrill.

We have included an additional copy of the response as well as the copy of the sending unit's report confirming transmission.

Respectfully submitted,

Joan Simunic, Reg. No. 43,125

Sild-Chemie Inc.



United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/851,177	05/08/2001	X.D. Hu	ZL494/01001	2722
22884	7590 11/18/2003		EXAMI	NER
	ON & REUTLINGER N & WILLIAMSON TOWER		NGUYEN,	CAM N
	E, KY 40202		ART UNIT	PAPER NUMBER
			1754	
2 to Table 10 30 Hotel		•	DATE MAILED: 11/18/2003	dict
<u>\$</u>			The state of the second of	tom - A - C -

- Notice of abendonment

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PTO-90C (Rev. 10/03)

<u>.</u>	IPE			
·	Notice of Abandonment 1 Application No. 09/851,177	Applicant(s)	Hu et	af.
	Cam Nguye		Art Unit 1754	
	The MAILING DATE of this communication pears on the cover sheet with	h the corres	pondence addr	
This	application is abandoned in view of:	•		
1. 🛛	Applicant's failure to timely file a proper reply to the Office letter mailed	on <i>Mar 2i</i>	5 2003	
(a)	A reply was received on (with a Certificate of Ma), which is after the expiration of the period for re month(s)) which expired on	ilina es Tes		d nsion of time of
(b)	A proposed reply was received on, but it does not 1.113(a) to the final rejection.			
	(A proper reply under 37 CFR 1.113 to a final rejection consists only of: (the application in condition for allowance; (2) a timely filed Notice of Application Continued Examination (RCE) in compliance with 37 CFR 1.1	ear (with ap 14).	peal fee); or (3) a timely filed
(c)	A reply was received on but it does not constitute proper reply, to the non-final rejection. See 37 CFR 1.85(a) and 1.111	a proper ro . (See expl	eply, or a bona	fide attempt at a
(d)	No reply has been received.			7 below).
2.	Applicant's failure to timely pay the required issue fee and publication fee three months from the mailing date of the Notice of Allowance (PTOL-85)	, if applicat	ole, within the	statutory period of
(a)	The issue fee and publication fee, if applicable, was received on	he statutor	(with a Certifi y period for pa	cate of Mailing or syment of the issue
(þ)	The submitted issue fee of \$ is insufficient. A balance of \$	is due	·) .	
	The issue fee required by 37 CFR 1.18 is \$ The publication fe	e, if require	ed by 37 CFR	1.18(d) is \$
(c)	☐ The issue fee and publication fee, if applicable, has not been received.			
3. 🗌	Applicant's failure to timely file corrected drawings as required by, and w Notice of Allowability (PTO-37).	ithin the th	ee-month peri	od set in, the
(a)	Proposed new formal drawings were received on(Transmission dated), which is after the expiration	with a Cert	ificate of Maili	ng or
(b)	☐ No corrected drawings have been received.	or the perio	od for reply.	•
4.	The letter of express abandonment which is signed by the attorney or againterest, or all of the applicants.	ent of record	d, the assigned	of the entire
5.□	The letter of express abandonment which is signed by an attorney or ager under 37 CFR 1.34(a)) upon the filing of a continuing application.	nt (acting in	a representat	ive capacity
6. 🗆	The decision by the Board of Patent Appeals and Interferences rendered operiod for seeking court review of the decision has expired and there are not become	n_ no allowed	and	d because the
7. 🗆	The reason(s) below:			

Petitions to revive under 37 CFR 1.137(a) or (b), or requests to withdraw the holding of abandonment under 37 CFR 1.181, should be promptly filed to minimize any negative effects on patent term.

TO:Auto-reply fax to 502 o34 7661 COMPANY:



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TO:

Fax Sender at 502 634 7661

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09/24/03 FED 07:47 PAX 502 634 7661

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U.S. Patent and Trademark Office Examiner: Cam Nguyen Gro Fax #703-872-9310 Croup: 1754

Joan L.Simunic Reg. No. 42,125

FROM: CATE:

September 24, 2003

PAGES .

in total (including cover sheet) O.S. Patent Application No.09/851,177

Remarks: This faculmile is in mesponed to the office action, having a mailing date of March 26, 2003

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 (2) Foo Transmittel for FV 2003
 (3) Ctedit Card Payment Form
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9/24/03 Cato

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U.S. Patent and Trademark Office

Examiner: Cam Nguyen

Group: 1754

Fax #703-872-9306

FROM:

Joan L. Simunic Reg. No. 43,125

DATE:

September 24, 2003

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in total (including cover sheet) 20

RE:

U.S. Patent Application No.09/851,177

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Donna Ferrill

Person Signing

Butter the Paperwork Reduction Act of 1995	U.S. Pat	PTO/SB/21 (08-03) Approved for use through 08/30/2003. OMB 0651-0031 tent and Trademark Office; U.S. DEPARTMENT OF COMMERCE tion of information unless it displays a valid OMB control number.		
TRANSMITTAL	Application Number	09/851,177		
TRANSMITTAL	Filing Date	May 8, 2001		
FORM	First Named Inventor	X. D. Hu		
(to be used for all correspondence after initial fil	ng) Art Unit	1754		
	Examiner Name			
	O Attorney Docket Number	Cam Nguyen		
Total Number of Pages in This Submission	,	ZL 494/01001		
	ENCLOSURES (Check all th	at apply)		
Fee Transmittal Form Drawing(s)				
Response to Missing Parts under 37 CFR 1.52 or 1.53	IDE OF ADDI ICANT ATTORN			
Firm Joan L Simunic Reg. 43 13	JRE OF APPLICANT, ATTORN	IET, UK AGENT		
or Individual name Signature Date September 23, 2008	of and a second			
CES	TIEICATE OF TRANSMISSION			
I hereby certify that this correspondence is beir sufficient postage as first class mail in an envel the date shown below. Typed or printed name	RTIFICATE OF TRANSMISSION og facsimile transmitted to the USPTO of ope addressed to: Commissioner for Pa	N/MAILING r deposited with the United States Postal Service with atents, P.O. Box 1450, Alexandria, VA 22313-1450 on		
Signature // WWW.	full	Date Whowher 24,2003		

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for FY 2003	

Effective 01/01/2003. Patent fees are subje

Jean L. Simunic

Name (Print/Type)

Signature

Applicant claims small entity status. See 37 CFR 1.27

(\$) 930.00 TOTAL AMOUNT OF PAYMENT

Complete if Known		
Application Number	09/851,177	
Filing Date	May 8, 2001	
First Named Inventor	X. D. Hu	
Examiner Name	Cam Nguyen	
Art Unit	1754	
Attorney Docket No.	ZL 494/01001	_

METHOD OF PAYMENT (check all that apply) FEE CALCULATION (continued)				
Check Credit card Money Other None	3. ADDITIONAL FEES			
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Deposit Account	1051 130 2051 65 Surcharge - late filing fee or oath	414		
Number Deposit	1052 50 2052 25 Surcharge - late provisional filing fee or -			
Account Name	cover sheet			
The Director is authorized to: (check all that apply)	1053 130 1053 130 Non-English specification 1812 2 520 1812 2 520 For filing a request for ex parte reexamination			
Charge fee(s) indicated below Credit any overpayments	4904 020* 1904 020* Requesting publication of SIR prior to			
Charge any additional fee(s) during the pendency of this application	n Examiner action			
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to the above-identified deposit account.	1251 110 2251 55 Extension for reply within first month			
FEE CALCULATION	1252 410 2252 205 Extension for reply within second month			
1. BASIC FILING FEE	1252 410 2232 203 1253 930 1253 465 Extension for reply within third month	0		
Large Entity Small Entity Fee Fee Fee Fee Fee Description Fee Paid	1254 1,450 2254 725 Extension for reply within fourth month			
Code (\$) Code (\$)	1255 1,970 2255 985 Extension for reply within fifth month			
1001 750 2001 375 Utility filing fee	1401 320 2401 160 Notice of Appeal			
1002 330 2002 165 Design filing fee	1401 320 2401 100 Notice of Appeal			
1003 520 2003 260 Plant filing fee	1403 280 2403 140 Request for oral hearing			
1004 750 2004 375 Reissue filing fee	1451 1.510 1451 1.510 Petition to institute a public use proceeding			
1005 160 2005 80 Provisional filing fee	1452 110 2452 55 Petition to revive - unavoidable			
SUBTOTAL (1) (\$) n/a	1453 1.300 2453 650 Petition to revive - unintentional			
2. EXTRA CLAIM FEES FOR UTILITY AND REISSUE	E 1501 1.300 2501 650 Utility issue fee (or reissue)			
Fee from Extra Claims below Fee Paid				
Total Claims X =	1503 630 2503 315 Plant issue fee			
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Multiple Dependent =	1807 50 1807 50 Processing fee under 37 CFR 1.17(q)			
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Fee Fee Fee Fee Fee Description Code (\$)	8021 40 8021 40 Recording each patent assignment per			
1202 18 2202 9 Claims in excess of 20	property (times number of properties)			
1201 84 2201 42 Independent claims in excess of 3	1809 750 2809 375 Filing a submission after final rejection (37 CFR 1.129(a))			
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**or number previously paid, if greater; For Reissues, see above	*Reduced by Basic Filing Fee Paid SUBTOTAL (3) (\$) 930			
SUBMITTED BY	(Complete (# applicable))			

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Registration No.

(Attorney/Agent)

43,125

Telephone 502-634-7373

Date

September 23, 2003

This collection of information is required by 37 CFR 1.17 and 1.27. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

PTO/SB/22 (08403)

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PETITION FOR EXTENSION OF TIME UNDER 37 CFR 1.136(a)			Docket Number (Optional) 71494/0100 (
			D. Hu et al	
ł		Application Number 09	1851, 177 Filed May 8, 2001	
	For High Surface Agea			
		Art Unit 1754	Examiner C. Mauyen	
This is a request under the provisions of 37 CFR 1.136(a) to extend the period for filing a reply in the above identified application.				
The requested extension and appropriate non-small-entity fee are as follows (check time period desired):				
•	One month (37 CFR 1.17(a)(1))		\$	
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	Four months (37 CFR 1.17(a)(4))		\$	
	Five months (37 CFR 1.17(a)(5))		\$	
	Applicant claims small entity status. See 37 CFR 1.27. Therefore, the fee amount shown above is reduced by one-half, and the resulting fee is: \$			
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⊠ ⊠	Payment by credit card. Form PTO-2038 is attached. The Director has already been authorized to charge fees in this application to a Deposit Account.			
	The Director is hereby authorized to che to Deposit Account Number	Director is hereby authorized to charge any fees which may be required, or credit any overpayment, eposit Account Number		
	have enclosed a duplicate copy of this sheet.			
	I am the applicant/inventor.			
	assignee of record of the entire interest. See 37 CFR 3.71. Statement under 37 CFR 3.73(b) is enclosed (Form PTO/SB/96). attorney or agent of record. Registration Number 43, 125			
	attorney or agent unde Registration number if a	er 37 CFR 1.34(a). cting under 37 CFR 1.34(a)		
	WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.			
	September 23, 200		Mence	
	Date Signature			
	Telephone Number Joan L. Simunic Typed or printed name			
NOTE: Signatures of all the inventors or assignees of record of the entire interest or their representative(s) are required. Submit multiple forms if more than one signature is required, see below.				
Total of forms are submitted.				

This collection of information is required by 37 CFR 1.136(a). The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 6 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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PETITION FOR EXTENSION OF TIME UNDER 37 CFR 1.136(a) Docket Number (Optional) 71494/0100 1			
In re Application of X.D. Hu et al			
Application Number 09/851, 177 Filed May 8, 2001			
For High Surface Acea			
Art Unit 1754 Examiner C. Navyen			
This is a reconstitution the provisions of 37 CFR 1.136(a) to extend the period for filing a reply in the above identified application.			
The requested extension and appropriate non-small-entity fee are as follows (check time period desired):			
One month (37 CFR 1.17(a)(1))			
Two months (37 CFR 1.17(a)(2))			
Three months (37 CFR 1.17(a)(3)) \$ 930			
Four months (37 CFR 1.17(a)(4))			
Five months (37 CFR 1.17(a)(5))			
Applicant claims small entity status. See 37 CFR 1.27. Therefore, the fee amount shown above is reduced by one-half, and the resulting fee is: \$			
A check in the amount of the fee is enclosed.			
ayment by credit card. Form PTO-2038 is attached.			
ne Director has already been authorized to charge fees in this application to a Deposit Account.			
he Director is hereby authorized to charge any fees which may be required, or credit any overpayment, Deposit Account Number			
I have enclosed a duplicate copy of this sheet.			
I am the applicant/inventor.			
assignee of record of the entire interest. See 37 CFR 3.71. Statement under 37 CFR 3.73(b) is enclosed (Form PTO/SB/96).			
attorney or agent of record. Registration Number 43, 125			
attorney or agent under 37 CFR 1.34(a). Registration number if acting under 37 CFR 1.34(a)			
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Syptember 23, 2003 Joan Signature			
Telephone Number Joan L. Simunic Typed or printed name			
NOTE: Signatures of all the inventors or assignees of record of the entire interest or their representative(s) are required. Submit multiple forms if more than one signature is required, see below.			
Total of forms are submitted.			

This collection of information is required by 37 CFR 1.136(a). The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 6 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application Number

09/851,177

Applicants

X. D. Hu et al.

Filing Date

05/08/2001

TC/A.U.

1754

Examiner

Cam Nguyen

Attorney Docket No.

ZL494/01001

Customer No.

Title:

High Surface Area, Small Crystallite Size Catalyst for Fischer-Tropsch

Synthesis

Commissioner for Patents P.O. Box 1450 Alexandria VA 22313-1450

ADMENDMENT

Sir:

In response to the Office Action mailed March 26, 2003, please amend the above-identified application as follows:

Amendments to the Specification: There are no changes are being made to the specification.

Amendments to the Claims are reflected in the listing of claims which begins on page 2 of this paper.

Amendments to the Drawings: There are no changes are being made to the drawings.

Remarks/Arguments begin on page 6 of this paper.

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1. (currently amended) A catalyst for use in the Fischer-Tropsch process, said catalyst comprising a catalyst particle, which comprises at least one metal that is an efficient carbon monoxide adsorber and at least one promoter, said metal and said promoter being dispersed on a support to form a said catalyst particle, said particle having a BET surface area of from about 100 m²/g to about 250 m²/g, and said metal and said promoter being are dispersed on the support such that the crystallite size of the metal oxide is from about 40 Å to about 200 Å, and said particle having an essentially smooth, homogeneous surface morphology.

Claim 2. (original) The catalyst of Claim 1 wherein said particle comprises from about 5 wt % to about 60 wt % cobalt, and from about 0.0001 wt % to about 1 wt % of a first promoter, and from about 0.01 wt % to about 5 wt % of a second promoter.

Claim 3. (original) The catalyst of Claim 2 wherein said particle comprises from about 10 wt% to about 30 wt % cobalt, and from about 0.01 wt % to about 0.05 wt % of said first promoter, and from about 0.1 wt % to about 1 wt % of said second promoter.

Claim 4. (original) The catalyst of Claim 1 wherein said metal is selected from the group consisting of nickel, cobalt, iron, ruthenium, osmium, platinum, palladium, iridium, rhenium, molybdenum, chromium, tungsten, vanadium, rhodium, copper, zinc, and combinations thereof.

Claim 5. (original) The catalyst of Claim 4 wherein said metal is cobalt.

Claim 6. (currently amended) The catalyst of Claim 1 wherein said promoter is selected from the group consisting of boron, cerium, chromium, copper, iridium, iron, lanthanum, manganese, molybdenum, palladium, platinum, rhenium, rhodium, ruthenium, strontium, tungsten, vanadium, zinc, sodium oxide, potassium oxide, rubidium oxide, cesium oxide, magnesium oxide, titanium oxide, zirconium oxide, and other rare earth metals, such as scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, other rare earth metals and combinations thereof.

Claim 7. (original) The catalyst of Claim 2 wherein said first promoter is selected from the group consisting of palladium, platinum, ruthenium, rhenium, rhodium, iridium and a combination thereof; and said second promoter is selected from the group consisting of potassium, boron, cesium, lanthanum, cerium, strontium, scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, palladium, platinum, ruthenium, rhenium, rhodium, iridium and combinations thereof.

Claim 8. (original) The catalyst of Claim 1 wherein said support is selected from the group consisting of aluminum oxide, γ-alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silica, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.

Claim 9. (original) The catalyst of Claim 8 wherein said support is γ -alumina.

- Claim 10. (original) The catalyst of Claim 9 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.
- Claim 11. (currently amended) A catalyst for use in the Fischer-Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size, and wherein said particle being is formed by the steps of:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- c) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is essentially completely reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles; and
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours.
- Claim 12. (original) The catalyst of Claim 11 wherein said support is selected from the group consisting of aluminum oxide, γ-alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silica, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
 - Claim 13. (original) The catalyst of Claim 12 wherein said support is aluminum oxide.
- Claim 14. (original) The catalyst of Claim 12 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.
- Claim 15. (original) The catalyst of Claim 11 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.
- Claim 16. (currently amended) The catalyst of Claim 15 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water molecules, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.
- Claim 17. (original) The catalyst of Claim 16 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate.
- Claim 18. (original) The catalyst of Claim 11 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).

- Claim 19. (original) The catalyst of Claim 11 wherein said solid portion is dried at from about 120°C to about 260°C in step f).
- Claim 20. (original) The catalyst of Claim 11 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- Claim 21. (original) The catalyst of Claim 11 wherein said particles are further stabilized to prevent pyrophoric reactions when said particles are in the presence of air.
- Claim 22. (currently amended) The catalyst of Claim 21 wherein said particles are stabilized by being coated coating with oil.
- Claim 23. (currently amended) The catalyst of Claim 11 further including at least one promoter, wherein said promoter being is added with said cobalt salt solution.
- Claim 24. (original) The catalyst of Claim 23 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 25. (original) The catalyst of Claim 11 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.

Claims 26 – 34. (canceled)

- 26. A method for making a catalyst for use in the Fischer Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle, said method comprising:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b)——adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 120°C;
- e) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is essentially completely reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles; and
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours.
 - 27. The catalyst of Claim 26 wherein said support is aluminum oxide.
- 28. The catalyst of Claim 27 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate.

- 29. The catalyst of Claim 26 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).
 - 30. The catalyst of Claim 26 wherein said solid portion is dried at from about 120°C to about 260°C in step f).
- 31. The catalyst of Claim 26 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- 32. The catalyst of Claim 26 wherein said particles are further stabilized by being coated with oil.
- 33. The catalyst of Claim 26 further including at least one promoter, said promoter being added with said cobalt salt solution.
- 34.— The catalyst of Claim 33 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 35. (new) A catalyst for use in the Fischer-Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle, wherein said particle is formed by the steps of:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- c) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles;
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours; and
- h) stabilizing said particles to prevent pyrophoric reactions when said particles are in the presence of air by coating said particles with oil.
- Claim 36. (new) The catalyst of Claim 35 wherein said support is selected from the group consisting of aluminum oxide, γ-alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silicate, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
- Claim 37. (new) The catalyst of Claim 36 wherein said support has a particle size of from about 60 μm to about 150 μm, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.

- Claim 38. (new) The catalyst of Claim 35 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.
- Claim 39. (new) The catalyst of Claim 38 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.
- Claim 40. (new) The catalyst of Claim 35 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).
- Claim 41. (new) The catalyst of Claim 35 wherein said solid portion is dried at from about 120°C to about 260°C in step f).
- Claim 42. (new) The catalyst of Claim 35 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- <u>Claim 43. (new)</u> The catalyst of Claim 35 further including at least one promoter wherein said promoter is added with said cobalt salt solution.
- Claim 44. (new) The catalyst of Claim 43 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 45. (new) The catalyst of Claim 35 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.
- Claim 46. (new) The catalyst of Claim 35 wherein said support is aluminum oxide and said aqueous cobalt salt solution comprises water and hexaammine cobalt (II) carbonate.

Attachment: Clean Unmarked Version of Claims Now in Application

REMARKS / ARGUMENTS

Remarks Regarding Informalities and Claims Rejected Under 35 USC §112

Claims 1-25 remain in the application. Claims 35-46 have been added to the application and are presented for examination. Claims 26-34 have been withdrawn in response to a requirement by the Examiner that the claims be restricted to Claims 1-25, drawn to a catalyst, or to Claims 26-34, drawn to a process of preparing a catalyst. In view of the Examiner's earlier restriction requirement, applicant retains the right to present claims 26-34 in a divisional application.

The Examiner objected to claims 1, 11, 16, 22, and 23 because of noted informalities. As proposed by the Examiner, claims 1 and 11 have been reworded in lines 1 – 3. Further, as proposed by the Examiner: in claim 1, line 5, "being" has been replaced with "are"; in claim 11, line 2, "being" has been replaced with "is"; and, in claim 16, line 2, "molecules" has been deleted to retain consistency with the language of claim 15. Further formality amendments include: in claim 22, line 1, "being coated" has been deleted and replaced with "coating"; and in claim 23, line 2, "being" has been replaced with "is" and "wherein" has been added. The amendments of claim 22 and claim 23 were suggested by the Examiner but required further correction than proposed to have grammatically correct sentences after amendment. Finally, in claim 11, step (c), "essentially completely" has been deleted to address a concern raised by the Examiner with the original language. No new subject matter has been added in making these amendments.

The Examiner objected to claim 17 based on the failure to comply to 37 CFR 1.75(c). The applicant respectfully challenges this objection. Claim 17 depends from claim 16 which recites a cobalt (II) complex having "coordination sphere ligands selected from the group consisting of water, ammonia, ..." Claim 17 recites "The catalyst of Claim 16 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate." As is known in the art, the "hexaammine cobalt (II) carbonate" coordination sphere ligands are ammonia, as recited in claim 16. The carbonate is not a coordination sphere ligand but rather a counterion to the positively charged cobalt ion. Thus, applicant requests the Examiner to reconsider his objection with respect to claim 17.

Claims 1 – 10 and 16 – 17 were rejected under 35 U.S.C. §112. The Examiner states that Claim 1 recites the limitation "the metal oxide" with insufficient antecedent basis for this limitation. The applicant respectfully notes that in the specification on page 4, lines 2 – 3, the catalyst has "a metal oxide crystallite size of from about 40 Å to about 200 Å." On page 12, lines 11 – 15, the metal oxide is more narrowly defined as the "cobalt oxide crystallite size" – and as is known in the art, cobalt is considered a metal when in reference to oxides – and the "cobalt oxide crystallite size ... is greater than about 40 Å diameter, and is preferably less than about 200 Å." Thus, applicant requests the Examiner to reconsider his objection under 35 U.S.C. §112 with respect to claim 1, and its dependent claims 2 – 10. If it would expedite matters, the applicant is willing to amend the paragraph on page 12 to include the more inclusive language of page 4 if so requested by the Examiner.

As noted by the Examiner, Claim 6 included the indefinite phrase "such as". Claim 6 has been amended to delete this phrase and should now be in allowable form.

As noted by the Examiner, Claim 16 included improper Markush terminology. Claim 16 has been amended to be consistent with the correct Markush phrase, and claim 16 and its dependent claim 17 should now be in allowable form.

Remarks Regarding Allowable Subject Matter

The Examiner has indicated that Claim 22 is objected to as being dependent on a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims. In response to this rejection, new claim 35 is presented. New claim 35 includes the limitations of rejected independent claims 11 (preamble and steps a-g), rejected dependent claim 21 and objected to dependent claim 22 (step h). Claims 36-46 have been added as claims dependent on new independent claim 35. As amended, independent claim 35 is believed to now be in allowable form. Because they depend from an allowable claim, dependent claims 36-46 are also believed to now be in allowable form.

The Examiner has also indicated that claims 1-10 would be allowable if the 35 U.S.C. §112 rejection is overcome. As noted in the prior section, page 4, lines 2-3, of the specification teach a catalyst that has "a metal oxide crystallite size of from about 40 Å to about 200 Å" and page 12, lines 11 -15, of the specification teach "cobalt oxide crystallite size ... greater than about 40 Å diameter, and is preferably less than about 200 Å." Applicant is willing to amend the paragraph on page 12 to include the more inclusive language of page 4 if so requested by the Examiner.

Remarks Regarding Citations

Applicant has made note of the prior art recited by the Examiner in Paragraph 15 of his Office Action mailed March 26, 2003.

Remarks Regarding Claims Rejected Under 35 USC §102(b) and 35 USC §103(a)

The Examiner has rejected independent claim 11 and dependent claims 12, 14 – 21, 23 and 25 under 35 U.S.C 102(b) as anticipated by or, in the alternative, under 35 U.S.C. 103(a) as obvious over Shen et al, U.S. Patent 5,962,367 ("the '367 patent"). Applicant respectfully traverses both rejections.

The Examiner has further rejected independent claim 11 and dependent Claims 12-13, 15-21 and 23-25 under 35 U.S.C 102(b) as anticipated by or in the alternative, under 35 U.S.C. 103(a) as obvious over Sapienza et al., U.S. Patent 4,396,539 ("the '539 patent"). Applicant respectfully traverses both rejections.

Summary of the Present Invention

The Fischer-Tropsch catalyst of the present invention is a transition metal-based catalyst having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support, and a small metal oxide crystallite size. In a first embodiment, the catalyst has a surface area of from about 100 m²/g to about 250 m²/g; an essentially smooth, homogeneous surface morphology; an essentially uniform distribution of metal throughout an essentially inert support; and a metal oxide crystallite size of from about 40 Å to about 200 Å. Optionally, the catalyst may further comprise at least one promoter.

The Fischer-Tropsch catalyst of independent claim 11 of the present application, as amended, is a catalyst comprising cobalt dispersed on a support to form a catalyst particle having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size, and wherein the particle is formed by several prescribed steps. Claims 12 – 25 of the present application depend from independent claim 11.

As provided in dependent claim 14, the catalyst of claim 11 may be prepared from a support having a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm. However, as discussed on page 28, line 1 through page 29, line 4 and as shown in Figures 1A – 4, two catalysts prepared with the same support material can have significantly different physical characteristics depending upon the process used to make the catalyst. (N.B.: In the examples of the present application each catalyst was prepared using a Puralox SCCa 5/150 support having a surface area of about 160 m²/g, a pore volume of about 0.50 ml/g, and a pore diameter of about 12.55 nm.) The steps outlined in independent claim 11 result in a catalyst particle having the desired physical characteristics. Prior art methods of catalyst particle preparation, even starting with the same support, do not result in the catalyst particle of the present invention.

Summary of U.S. Patent 5,962,367, Shen et al.

U.S. Patent 5,962,367 teaches a process for preparing a catalyst support primarily formed of titania. The catalyst support comprises 60 wt% to 100 wt % titania as TiO_2 and 0 wt% to 40 wt % alumina as Al_2O_3 . As noted by the Examiner, the '367 patent teaches a cobalt molybdate catalyst supported on a titania support, wherein the support has a surface area ranging from 80 m²/g to 200 m²/g, a pore volume of from about 0.3 ml/g to about 0.5 ml/g, and a pore diameter of from about 6 nm to about 20 nm.

The '367 patent then teaches using this catalyst support to prepare a cobalt molybdate catalyst using an impregnation process. At column 5, lines 6 – 8, the '367 states "the dynamic co-impregnation

method is used for supporting the cobalt and molybdenum active components ..." In Examples 4-7, the support is "immersed into the impregnating solution ..." for the preparation of the catalyst.

However, the '367 patent does not teach or suggest that the catalyst resulting from the impregnation process using the '367 titania support has a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size as is seen with the preparation process of the present invention. Rather, there is nothing in the '367 patent that suggests that the physical characteristics of the cobalt molybdate catalyst prepared by the impregnation process would be any different than the catalysts prepared by the prior art impregnation method and reported in Example 6 (and Figures 2A, 2B and 4) of the present application. Further, the catalyst of the present invention made by the process of instant claim 11 would not be anticipated or obvious merely because a titania support is used, or the physical characteristics of the support are similar to supports of the prior art, or because a promoter is added. Thus, independent claim 11 and its dependent claims 12 – 25 are not anticipated nor obvious in view of U.S. Patent 5,962,367.

Summary of U.S. Patent 4,396,539, Sapienza et al.

U.S. Patent 4,396,539 teaches a process for preparing a catalyst suitable for Fischer-Tropsch synthesis wherein the catalyst is composed of palladium or platinum and cobalt supported on a solid support. More specifically, the '539 patent teaches an alumina, silica gel, kieselguhr, or zinc oxide support that is impregnated by immersion of the support in an aqueous solution of the salt of a palladium or platinum metal. The palladium- or platinum-treated support is then immersed in a solution containing primarily cobalt.

The objective of the process set forth in the '539 patent is to achieve a catalyst particle with the metal loading as a coating around the support. The catalyst structure depicted in Figure 5 is believed to "best represent the idealized structure of this catalyst" (column 4, lines 9-21). By contrast, the process of claim 11 of the present invention produces a catalyst that has "an essentially uniform distribution of cobalt throughout the support". In other words, the process taught in the '539 patent teaches away from the desired product of the present invention. Thus, independent claim 11 and its dependent claims 12-25 are not anticipated nor obvious in view of U.S. Patent 4,396,539.

Because U.S. Patent 5,962,367 and U.S. Patent 4,396,539 each teach a process for preparing a catalyst that relies on impregnation to deliver the metal to the support, the product-by-process taught by the amended claim 11 of the present application is neither anticipated nor obvious. Thus, applicant respectfully requests that the $\S102(b)$ and the $\S103(a)$ rejections be withdrawn and that independent claim 11, as amended, and its dependent claims 12-25, be allowed.

Applicant respectfully requests that a timely Notice of Allowance be issued in this case.

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Clean Unmarked Version of Claims Now in Application

- Claim 1. A catalyst for use in the Fischer-Tropsch process, said catalyst comprising a catalyst particle, which comprises at least one metal that is an efficient carbon monoxide adsorber and at least one promoter dispersed on a support to form said catalyst particle, said particle having a BET surface area of from about $100 \text{ m}^2/\text{g}$ to about $250 \text{ m}^2/\text{g}$, and said metal and said promoter are dispersed on the support such that the crystallite size of the metal oxide is from about 40 Å to about 200 Å, and said particle having an essentially smooth, homogeneous surface morphology.
- Claim 2. The catalyst of Claim 1 wherein said particle comprises from about 5 wt % to about 60 wt % cobalt, and from about 0.0001 wt % to about 1 wt % of a first promoter, and from about 0.01 wt % to about 5 wt % of a second promoter.
- Claim 3. The catalyst of Claim 2 wherein said particle comprises from about 10 wt% to about 30 wt % cobalt, and from about 0.01 wt % to about 0.05 wt % of said first promoter, and from about 0.1 wt % to about 1 wt % of said second promoter.
- Claim 4. The catalyst of Claim 1 wherein said metal is selected from the group consisting of nickel, cobalt, iron, ruthenium, osmium, platinum, palladium, iridium, rhenium, molybdenum, chromium, tungsten, vanadium, rhodium, copper, zinc, and combinations thereof.
 - Claim 5. The catalyst of Claim 4 wherein said metal is cobalt.
- Claim 6. The catalyst of Claim 1 wherein said promoter is selected from the group consisting of boron, cerium, chromium, copper, iridium, iron, lanthanum, manganese, molybdenum, palladium, platinum, rhenium, rhodium, ruthenium, strontium, tungsten, vanadium, zinc, sodium oxide, potassium oxide, rubidium oxide, cesium oxide, magnesium oxide, titanium oxide, zirconium oxide, scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, other rare earth metals and combinations thereof.
- Claim 7. The catalyst of Claim 2 wherein said first promoter is selected from the group consisting of palladium, platinum, ruthenium, rhenium, rhodium, iridium and a combination thereof; and said second promoter is selected from the group consisting of potassium, boron, cesium, lanthanum, cerium, strontium, scandium, yttrium, praseodymium, neodymium, promethium, samarium, europium, gadolinium, terbium, dysprosium, holmium, erbium, thulium, ytterbium, lutetium, palladium, platinum, ruthenium, rhenium, rhodium, iridium and combinations thereof.
- Claim 8. The catalyst of Claim 1 wherein said support is selected from the group consisting of aluminum oxide, γ-alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silica, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
 - Claim 9. The catalyst of Claim 8 wherein said support is γ -alumina.
- Claim 10. The catalyst of Claim 9 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.

- Claim 11. A catalyst for use in the Fischer-Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle_having a high surface area, a smooth, homogeneous surface morphology, an essentially uniform distribution of cobalt throughout the support and a small metal oxide crystallite size, and wherein_said particle is formed by the steps of:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- c) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles; and
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours.
- Claim 12. The catalyst of Claim 11 wherein said support is selected from the group consisting of aluminum oxide, γ -alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silica, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
 - Claim 13. The catalyst of Claim 12 wherein said support is aluminum oxide.
- Claim 14. The catalyst of Claim 12 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.
- Claim 15. The catalyst of Claim 11 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.
- Claim 16. The catalyst of Claim 15 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.
- Claim 17. The catalyst of Claim 16 wherein said cobalt (II) complex is hexaammine cobalt (II) carbonate.
- Claim 18. The catalyst of Claim 11 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).
- Claim 19. The catalyst of Claim 11 wherein said solid portion is dried at from about 120°C to about 260°C in step f).

- Claim 20. The catalyst of Claim 11 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- Claim 21. The catalyst of Claim 11 wherein said particles are further stabilized to prevent pyrophoric reactions when said particles are in the presence of air.
 - Claim 22. The catalyst of Claim 21 wherein said particles are stabilized by coating with oil.
- Claim 23. The catalyst of Claim 11 further including at least one promoter wherein said promoter is added with said cobalt salt solution.
- Claim 24. The catalyst of Claim 23 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 25. The catalyst of Claim 11 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.
- Claim 35. A catalyst for use in the Fischer-Tropsch process, said catalyst comprising cobalt dispersed on a support to form a catalyst particle, wherein said particle is formed by the steps of:
- a) adding said support to water, with agitation, to form a slurry, and maintaining a slurry temperature at from about 35°C to about 210°C;
- b) adding an aqueous cobalt salt solution having a pH value greater than the point of zero charge of said support to said slurry with agitation and while maintaining said slurry temperature at from about 65°C to about 120°C;
- c) agitating said slurry and maintaining said slurry temperature at from about 65°C to about 120°C until said cobalt salt is reacted with said support;
 - d) separating said slurry into a solid portion and a liquor portion;
 - e) washing said solid portion with water;
- f) drying and calcining said solid portion at from about 90°C to about 375°C to form catalyst particles;
- g) reducing said catalyst particles by heating said particles from ambient temperature to from about 300°C to about 500°C at a rate of from about 0.1°C/min to about 10°C/min over a period of from about 5 hours to about 40 hours; and
- h) stabilizing said particles to prevent pyrophoric reactions when said particles are in the presence of air by coating said particles with oil.
- Claim 36. The catalyst of Claim 35 wherein said support is selected from the group consisting of aluminum oxide, γ -alumina, alumina monohydrate, alumina trihydrate, alumina-silica, magnesium silicate, silica, silicate, silicalite, y-zeolite, mordenite, titania, thoria, zirconia, niobia, hydrotalcite, kieselguhr, attapulgite clay, zinc oxide, other clays, other zeolites and combinations thereof.
- Claim 37. The catalyst of Claim 36 wherein said support has a particle size of from about 60 μ m to about 150 μ m, a surface area of from about 90 m²/g to about 210 m²/g, a pore volume of from about 0.35 ml/g to about 0.50 ml/g, and a pore diameter of from about 8 nm to about 20 nm.

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- Claim 38. The catalyst of Claim 35 wherein said cobalt salt solution comprises water and a cobalt (II) complex having coordination sphere ligands selected from the group consisting of water, chloride ion, ammonia, pyridine, triphenylphosphine, 1,2-diaminoethane, diethylenetriamine, triethylenetetraamine, acetate, oxalate, 2,4-pentanedione, ethylenedinitilo tetraacetic acid, and combinations thereof.
- Claim 39. The catalyst of Claim 38 wherein said cobalt (II) complex has coordination sphere ligands selected from the group consisting of water, ammonia, pyridine, diaminoethane, diethylenetriamine, triethylenetetraamine, and a combination thereof.
- Claim 40. The catalyst of Claim 35 wherein the slurry temperature is maintained at from about 65°C to about 120°C in step a).
- Claim 41. The catalyst of Claim 35 wherein said solid portion is dried at from about 120°C to about 260°C in step f).
- Claim 42. The catalyst of Claim 35 wherein said particles are reduced in step g) by heating said particles from ambient temperature to about 350°C at a rate of about 1.0°C/min and then holding said particles at about 350°C for from about 12 hours to about 16 hours.
- Claim 43. The catalyst of Claim 35 further including at least one promoter wherein said promoter is added with said cobalt salt solution.
- Claim 44. The catalyst of Claim 43 wherein said promoter is a metal salt selected from the group consisting of rhenium (VII) oxide, ruthenium nitrosyl nitrate, platinum chloride, platinum ammine nitrate, platinum ammine chloride, and combinations thereof.
- Claim 45. The catalyst of Claim 35 further including at least one promoter impregnated onto said catalyst particle after said particle is dried in step f), said promoter being impregnated onto said particle by dipping said particle in an aqueous solution of said promoter while maintaining agitation, and then separating said impregnated particles from said solution, and drying said impregnated particles.
- Claim 46. The catalyst of Claim 35 wherein said support is aluminum oxide and said aqueous cobalt salt solution comprises water and hexaammine cobalt (II) carbonate.